FDA 510(k) Premarket Notification

K092073 MONOPOLY PEDICLE SCREW SYSTEM

CLEARED

Aug 7, 2009
30 days to decision

K092073 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k092073/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 8, 2009
Decision date	Aug 7, 2009
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Signus Medical**
Location	Washington, DC, US
Contact	THOMAS HOGHAUG
510(k) history	2 submissions · 2 cleared · 2009-2009